Attorney Docket No.: 8920-000005

## AMENDMENTS TO THE CLAIMS

The following listing of claims will replace all prior versions and listings of claims in the application.

## Listing Of Claims

- (Cancelled)
- (Cancelled) 2.
- (Currently Amended) A method for producing an enzyme collobiase, in the presence of glycoslyation inhibitor 2 deaxy D glucose from cultures of Termitomyces obspectus, said preparation contaming high concentration of enzyme collobiase in comparison to a control oulture, grown in absence of glycuslyation inhibitor 2 decay D glucoso, the said method comprising the steps of:
  - inoculating a mycelial culture of the Termitomyces clypeatus into sterilized medium containing carbon and nitrogen sources, inorganic salts, organic nutrients and glycosylation inhibitor 2-deoxy-D-glucose in the range of about 10 pg/ml 0.05 mg/ml to about 2 1 mg/ml at a pH of between about 3 to 8;
  - growing the mycelial culture at temperatures between 20-37°C under shaking acrobic conditions; and
  - separating culture medium from the mycelia to obtain the enzyme (c) preparation containing cellobiase activity[[,]] suid onzyme having an increased enzymatic activity in the range of about 1.15-2.236 units/ml to about 140.60 97 units/ml in the presence of glycosylation inhibiter 2 deaxy D glacose in comparison to cellobiase activity produced by the same organism under the same conditions in absence of the glycosylation inhibitor 2 deoxy D. gluceme.
  - (Cancelled) 4.

Attorney Docket No.: 8920-000005

- 5. (Cancelled)
- 6. (Cancelled)
- 7. (Currently Amended) The method as claimed in of claim 3, wherein the carbon source[[s]] of step (a) are is selected from the group consisting of carbohydrates, agrowastes, TCA cycle acids, amino acids, of and D-glucosamine, wherein the carbohydrates are selected from the group consisting of cellobiose, mannose, fructose, xylose, arabinose, starch, dextrine, cellulose, cotton, and xylan; wherein agrowastes are selected from the group consisting of baggasse powder, rice-straw powder, wheat bran, corn cob powder, and corn powder; wherein the TCA cycle acids are selected from the group consisting of succinate, fumarate, and maleate; and wherein the amino acids are selected from the group consisting of aspartate, glutamate, serine, histidine, and alanine.

HARNESS, DICKEY, & PIERCE

- 8. (Cancelled)
- 9. (Currently Amended) The method us obtained in of claim 3, wherein the nitrogen source in stop (a) is selected from the group consisting of ammonium chloride, ammonium nitrate, ammonium dihydrogen orthophosphate, and potassium nitrate.
- 10. (Currently Amended) The method as claimed in of claim 3, wherein the aterilized medium in step (a) comprises an organic nutrient selected from the group consisting of malt extract, yeast extract, potato extract, peptone, soya-peptone, bactopeptone, and com steep liquor.
- 11. (Currently Amended) The method acclaimed in of claim 3, wherein the sterilized medium further comprises a detergent selected from group consisting of Tween-20, Tween-80, and Tween-100.

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Attorney Docket No.: 8920-000005

## 12. (Cancelled)

- 13. (Currently Amended) The method as claimed in claim 8 of claim 3, wherein enhanced ensyme activity of celluliase is about 2.23 units/ml-in presence of about 0.05mg/ml of and the 2-deaxy-D-glucose is present at a concentration of about 0.05 mg/ml.
- 14. (Currently Amended) The method as claimed in claim 8 of claim 3, wherein enhanced enzyme activity of cellobiase is about 50.09 units/ml in presence of about 1 mg/ml of and the 2 deoxy-D-glucose is present at a concentration of about 1 mg/ml.
- 15. (Currently Amended) The method-as claimed in claim 8 of claim 3, wherein enhanced enzyme activity of collobiase is about 90 units/ml in presence of about 1 mg/ml of and the 2-deoxy-D-glucose is present at a concentration of about 300 μg/ml.
- 16. (Currently Amended) The method as claimed in claim 8 of claim 3, wherein contamed enzyme activity of cellobiase is about 140 units/ml, in presence of, about 500 μg/ml of the 2 deoxy-D-glucose is present at a concentration of about 1 mg/ml and mannose is present at a concentration of about 500 μg/ml.
  - 17. (Cancelled)
  - 18. (Cancelled)
  - 19. (now) A method for producing cellobiase, said method comprising:
  - (a) inoculating a mycelial culture of the *Termitomyces clypeatus* into a sterilized culture medium containing carbon and nitrogen sources, inorganic salts, organic nutrients at a pH of between 3 to 8 and a glycosylation inhibitor selected

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Attorney Docket No.: 8920-000005

from the group consisting of tunicamycin, 1-deoxynojirimycin, and gluconolactone;

- (b) growing the mycelial culture at temperatures between 20-37°C under shaking acrobic conditions; and
- (c) separating culture medium from the mycelia to obtain cellobiase activity in the range of about 1,2075 units/ml to about 6,1820 units/ml.
- 20. (new) The method of claim 19, wherein the glycosylation inhibitor is tunicarrycin at a concentration of 10 μg/ml and the cellobiase activity is about 1,2075 units/ml.
- 21. (new) The method of claim 19, wherein the glycosylation inhibitor is 1-deoxynojirimycin at a concentration of about 80 μM and the cellobiase activity is about 1.4085 units/ml.
- 22. (new) The method of claim 19, wherein the glycosylation inhibitor is glucono-lactone at a concentration of about 2 mg/ml and the cellobiase activity is about 6.1820 units/ml.
- 23. (new) The method of claim 3, wherein the pH is about 4.5 and the cellobiase activity is about 90 units/ml.
- 24. (new) The method of claim 3, wherein the pH is about 4.5, the carbon source is mannose, and the cellobiase activity is about 140 units/ml.
- 25. (new) The method of claim 7, wherein the carbon sources are selected from the group consisting of cellobiose, mannose and succinate.
- 26. (new) The method of claim 9, wherein the nitrogen source is ammonium dihydrogen orthophosphate.

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Attorney Docket No.: 8920-000005

- (new) The method of claim 10, wherein the sterilized medium comprises 27. potato extract.
- (new) The method of claim 3, wherein the Termitomyces chypeatus is a 28. Termitomyces clypeatus strain having Indian Institute of Chemical Biology accession number IICB-411.